-continued

	Cyclosporin	GLA	DGLA	EPA	DHA
F	0.05%	3%		1%	0.5%

I claim:

1. A dosage form of a fatty acid composition alone or in a pharmaceutical diluent or carrier comprising a cyclosporin and a fatty acid selected from the group consisting of GLA, DGLA and a derivative thereof convertible in the body thereto, wherein said fatty acid is in the range of 1 mg to 100 g, and wherein said cyclosporin is in the range of 0.1 ng to 100 mg per ml of said composition.

2. A dosage form of a composition according to claim 1 comprising also an n-3 essential fatty acid selected from the 20:5, 22:5 and 22:6 acids or their derivatives convertible in the body thereto.

3. A method of alleviating the renal side effects of administration of cyclosporin comprising administering a dosage form alone or in a pharmaceutical diluent or carrier to a person in need of same (1) a cyclosporin, and (2) a fatty acid selected from the group consisting of GLA, DGLA and a derivative thereof convertible in the body thereto, together or successively to a person suffering from or at risk of such side effects, wherein said fatty acid is in the range of 1 mg to 100 g, and wherein said cyclosporin is in the range of 0.1 ng to 100 mg per ml of said dosage form.

4. A dosage form of a composition according to claim 1 for administration of 50 mg to 10 g of the or each fatty acid and 250 mg to 1.5 g of the cyclosporin, per day.

5. A dosage form of a composition according to claim 1 comprising 0.1 to 10 mg per ml of the or each fatty acid and 0.1 ng to 1 mg per ml of the cyclosporin.

6. The method according to claim 3, wherein the GLA or DGLA is administered in an amount of 1 mg to 100 g daily and the amount of cyclosporin administered is 5 mg to 15 g daily.

7. The method according to claim 3, wherein the GLA or DGLA is administered in an amount of 50 mg to 10 g daily and the amount of cyclosporing administered is 250 mg to 1.5 g daily.

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